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L21 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2008 ACS on STN
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AN 2005:346977 CAPLUS

DN 142:392186

TI 2-Cyanobenzenesulfonamides for combating animal pests and their preparation

IN Von Deyn, Wolfgang; Baumann, Ernst; Hofmann, Michael; Kordes, Markus; Puhl, Michael; Schmidt, Thomas; Tedeschi, Livio; Rack, Michael; Bucci, Toni; Culbertson, Deborah L.; Cotter, Henry Van Tuyl; Oloumi-Sadeghi, Hassan

PA BASF Aktiengesellschaft, Germany

SO PCT Int. Appl., 67 pp.

CODEN: PIXXD2

DT Patent

FAN.CNT 1

LA English

L. LTIA .	AN.CHI I																		
	PATENT NO.			KIND DATE			APPLICATION NO.												
ΡI								WO 2004-EP11004											
		W :	AE.	AG.	AL.	AM.	AT.	AU,	AZ.	BA.	BB.	BG.	BR.	BW.	BY.	BZ.	CA.	CH.	
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	UΑ	2004		•		A1 20050421			AU 2004-279549					20041001					
		2539								CA 2004-2539563									
	ΕP	1670	752			A1		2006	0621		EP 2	004-	7657	61		20041001			
		R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,	
			IE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,	PL,	SK,	HR
	CN	1863	767			A		2006	1115	1	CN 2	004-	8002	8841		2	0041	001	
	BR	2004	0148	97		A		2006	1212		BR 2	004-	1489	7		20041001 20041001			
		2007						2007	0329		JP 2	006-	5300	74					
	MX	2006	PA03	145		Α		2006	0614	MX 2006-PA3145					20060320				
			A1		2007	0329	1						20060329 <						

20070817

20031002

20041001

IN 2006-CN1103

20060331

IN 2006CN01103

WO 2004-EP11004

PRAI US 2003-507507P

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W

The invention relates to 2-cyanobenzenesulfonamide compds. of formula I and/or to their agriculturally useful salts. In compds. I, R1 is C1-4 alkyl, C1-4 haloalkyl, C1-4 alkoxy, or C1-4 haloalkoxy; R2 is H, (un)substituted C1-6 alkyl, (un)substituted C2-6 alkenyl, (un)substituted C3-8 cycloalkyl, or (un)substituted C1-4 alkoxy; R3, R4, and R5 are independently selected from H, halo, cyano,

nitro, C1-6 alkyl, C3-8 cycloalkyl, C1-4 alkoxy, etc.; and includes agriculturally useful salts. The invention also relates to the preparation of I, agricultural compns. containing at least one compound of formula I and/or at least one agriculturally useful salt of I, and at least one inert liquid and/or solid agronomically acceptable carrier, and, if desired, at least one surfactant. Diazotization of 2-amino-6-methylbenzonitrile followed by sulfonylation and amidation with propylamine gave cyanobenzenesulfonamide II. Prepared similarly, the N-Et analog of II (I; R1 = Me; R2 = Et; R3 = R4 = R5 = H), at 300 ppm, resulted in over 75% mortality of 2-spotted spider mites (Tetranychus urticae) and over 85% mortality of 3 tested aphid species.

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

Uploading C:\Program Files\Stnexp\Queries\10574153-broad.str

Chain nodes : 8 9 10 11 ring nodes : 1 2 3 4 5 6 chain bonds : 6 -8 5-9 8-10 8-11 ring bonds : 1 -2 1 -6 2 -3 3-4 4-5 5-6 exact/norm bonds : 6 -6 8-9 8-10 8-11 normalized bonds : 1 -2 1 -6 2 -3 3-4 4-5 5-6 normalized bonds : 1-2 1-6 2-3 3-4 4-5 5-6 isolated ring systems : containing 1

Match level: 1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 8:CLASS 9:CLASS 10:CLASS 11:CLASS

STRUCTURE UPLOADED

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chain nodes : 1 2 3 4 5 6 7 8 chain bonds : 3-4 5-6 5-7 5-8

10574153

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FILE 'REGISTRY' ENTERED AT 12:58:14 ON 28 JAN 2008 SAV TEM L6 BRD574153/A

FILE 'REGISTRY' ENTERED AT 13:03:32 ON 28 JAN 2008 STRUCTURE UPLOADED 9 S L16 SAM SUB-L6 129 S L16 SSS FULL SUB-L6

FILE 'CAPLUS' ENTERED AT 13:03:58 ON 28 JAN 2008 16 S L18 1 S US200:-574153/APPS 1 S L19 AND L20 15 S L19 NOT L20

FILE 'REGISTRY' ENTERED AT 13:04:23 ON 28 JAN 2008

=> d 18 L8 HAS NO ANSWERS L8

Structure attributes must be viewed using STN Express query preparation.

Structure attributes must be viewed using STN Express query preparation.

-> d 122 tot bib abs hitstr

10574153 2 of 52 1-5 3-4 5-6 5-7 5-8 exact bonds : 1-2 1-3

Generic attributes : Saturation : Unsaturated

Number of Carbon Atoms : less than Type of Ring System : Monocyclic

STRUCTURE UPLOADED

Match level :

(FILE 'HOME' ENTERED AT 12:50:41 ON 28 JAN 2008)

FILE 'REGISTRY' ENTERED AT 12:50:51 ON 28 JAN 2008

SCREEN 1839

STRUCTURE UPLOADED

50 S (L3 AND L1) SAM

50 S L3 AND L1 NOT L2 SAM

55096 S L3 AND L1 NOT L2 SSS FULL L1 L2 L3 L4 L5 L6

FILE 'CAPLUS' ENTERED AT 12:53:02 ON 28 JAN 2008 : 57752 S L6 L7

FILE 'STNGUIDE' ENTERED AT 12:53:31 ON 28 JAN 2008

FILE 'REGISTRY' ENTERED AT 12:54:24 ON 28 JAN 2008 STRUCTURE UPLOADED L8 L9

26 S L8 SAM SUB=L6 611 S L8 SSS FULL SUB=L6 L10

FILE 'CAPLUS' ENTERED AT 12:54:55 ON 28 JAN 2008 280 S L10

FILE 'STNGUIDE' ENTERED AT 12:55:16 ON 28 JAN 2008

FILE 'REGISTRY' ENTERED AT 12:55:56 ON 28 JAN 2008 STRUCTURE UPLOADED 15 S L12 SAM SUB-L10 361 S L12 SSS FULL SUB-L10 L12 L13 L14

FILE 'CAPLUS' ENTERED AT 12:56:19 ON 28 JAN 2008 49 S L14 L15

FILE 'STNGUIDE' ENTERED AT 12:56:31 ON 28 JAN 2008 FILE 'CAPLUS' ENTERED AT 12:57:11 ON 28 JAN 2008

FILE 'STNGUIDE' ENTERED AT 12:57:24 ON 28 JAN 2008

10574153

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L22 ANSWER 1 OF 15 CAPLUS COPYRIGHT 2005 ACS on STN 2007:1145210 CAPLUS Full-text 147:421320

DN 147:421320
T1 3-Amino-1,2-benzisothiazole compounds for controlling invertebrate pests
TN Pohlman, Matchias; Von Deyn, Wolfgang; Kaiser, Florian; Baumann, Ernst;
Rack, Michael; Anspaugh, Douglas D.; Culbertson, Deborah L.; Van Tuyl
Cotter, Henry
PA BASF Aktiengesellschaft, Germany
SO PCT Int. Appl., 68pp.
CODEN: PIXXD2
DT Pater
LA English
FAN.CNT 1
FAN.CNT 1
FAN.CNT 1
FAN.CNT 1
FAN.CNT 10.
KIND DATE APPLICATION NO. DATE

KIND DATE APPLICATION NO. DATE

A1 20071011 MC 2007-EP\$2738 20070322

A1, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BM, BY, BZ, CA, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GH, GH, CH, LT, LT, LU, LY, MA, MD, MG, MK, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, SC, SD, SE, SG, SK, SL, SM, SY, SY, TJ, TM, TN, TR, TT, UG, US, UZ, VC, VN, ZA, ZM, ZM, BG, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, LT, LU, LY, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BP, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BP, CG, CI, CM, GA, GN, GQ, GM, ML, MR, NE, SN, TD, TG, BM, KE, LS, MM, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZM, AM, AZ, TT TM WO 2007113119

W: AE, AG,
CH, CN,
GD, GE,
KN, KP,
MN, MN,
RS, RU,
T2, UA, 

The present invention relates to pesticidal 3-amino-1,2-benzisothiazole compds. of formula [1], wherein Rl = (un)substituted (halo)alkyl, alkenyl, alkoxy, cycloalkyl, etc., R2-R4 = independently H, halo, CN, azido, NO2, (halo)alkyl, cycloalkyl, etc., R5 = H, OH, (un)substituted alkyl, etc., and n = 0, 1 or 2, or to enantiomers, disstereomers and salts thereof, with the provise that when n = 0, R5 is not H. The invention also relates compns. comprising such compds., application methods, and use of I derivs., their salts, or compns. comprising them for combating animal pests, including insects, arachnids or nematodes. Thus, cotton plants at the cotyledon stage were infested with cotton aphid (Aphis gossypii, mixed lite stages) prior to treatment with formulated solns, of I derivs., and aphid mortality was determined after 5 days. In this test 40 of the I derivs, at 300 pm provided >8st mortality of cotton ashid in comparison with untreseed controls. 286% mortality of cotton aphid in comparison with untreated controls. IТ

RL: RCT (Reactant); RACT (Reactant or reagent) (in preparation of aminobenzisothiazole pesticide) 850154-43-7 CAPLUS Benzenesulfonamide, 2-cyano-N-ethyl-3-methoxy- (CA INDEX NAME)

THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT RE.CNT 8

ANSWER 2 OF 15 CAPLUS COPYRIGHT 2008 ACS ON STN 2007:934877 CAPLUS <u>Full-text</u> 147:277460

Preparation of aminothiocarbonyl-substituted benzenesulfonamides and Preparation of aminocinocarbonyl-substituted benzenesuironasides and pyridinesulfonasides as pesticides for the protection of plants against animal pests and insects and particularly against aphids and thrips Kaiser, Florian; Von Deyn, Wolfgang, Pohlman, Matthias; Anspaugh, Douglas D.; Culbertson, Deborah L.; Cotter, Henry Van Tuyl
Basi Aktiengesellschaft, Germany IN

50 PCT Int. Appl., 259pp. CODEN: PIXXD2

DT Patent LA English FAN.CNT 1 PATENT NO.

P1 MO 200704530 A1 206/0823 WO 2007-EP51145 20070207
M: AE, AG, AL, AM, AT, AU, AZ, BA, BE, BG, BR, BM, BY, BZ, CA, CH,
CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, F1, GB, GD,
GE, GH, GM, GT, HN, HR, HU, ID, II, IN, IS, JP, KE, KG, MM, KY,
KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, KY,
KM, MM, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO,
RS, RU, SC, SD, SE, SG, SK, LS, SM, SV, SY, TJ, TM, TN, TR, TT,
TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZM
RN: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, F1, FR, GB, GR, HU, IE,
IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, 9E, S1, SK, TR, BF, BJ,
CF, CG, CT, CM, GA, GN, GO, GM, HM, NR, NB, SN, TD, TG, EM, GH,
GM, KE, LS, MM, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZM, AM, AZ, BY.

PRAI US 2006-774119P P 20060216
SI

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THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 3 OF 15 CAPLUS COPYRIGHT 2009 ACS ON STN 2007:595360 CAPLUS FJI1 text 147:3671

147:3671 Proparation of cyanobenzene derivatives as insecticides Ponlman, Matchias; Yon Deyn, Wolfgang; Schmidt, Thomas; Kaiser, Florian; Anspaugh, Douglas D.; Culbertson, Deborah L.; Van Tuyl Cotter, Henry Bast Aktiengesellschaft, Germany

PA SO PCT Int. Appl., 112pp. CODEN: PIXXD2

Patent English

FAN	CNT 1																
	PATENT	NO.			KIN	D	DATE			APPL	ICAT	ION	NO.		D.	ATE	
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PΙ	WO 200	70602	20		A2		2007	0531		WO 2	006-	EP68	880		2	0061	124
	WO 200	70502	20		A3		2007	0802									
	₩:	AE,	AG,	AL,	AM,	AΤ,	AU,	· AZ,	BA,	EB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH
		CN,	œ,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GE
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		KP,	KR,	KZ,	LA,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	LY.	MA,	MD,	MG,	MH
		MN,	MN,	MX,	MY,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RC
		RS,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SM,	sv,	SY,	TJ,	TM,	TN,	TR,	17
		TZ.	UA,	UG,	US,	UZ,	VC.	VN,	ZA,	ZM,	ZW						
	RM	: AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	ΗU,	ΙE
		IS,	IT,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI.	SK,	TR,	BF,	BJ
		CF,	α,	CI,	CM,	GΑ,	GN,	GQ,	G₩,	ML,	MR,	NE,	SN,	TD,	TG,	BW,	GH
		GM,	KE,	L9,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ ,	UG,	ZM,	ZW,	AM,	AZ,	BY
		KG,	ĸz.	MD,	RU,	TJ,	TM,	AP,	EA,	EP,	QΑ						
PRA	I US 200	5-739	642P		P		2005	1125									
os	MARPAT	147:	3671														
GI																	

The cyanobenzene derivs. I [m = 0, 1 or 2; A = N:CRSR6, N:SR7R3, NR10C(:X)R9 or ethylenically unsatd. or aromatic N-bound 5-, 6- or 7-membered heterocyclyl; X -0, S or NR11, R1 = H, nitro, cyano, saido, smino, halo,

The present invention relates to (thiotarbonyl)substituted aryl sulfonamides 1, wherein X is N, NO, CRS; R1 and R2 are independently H, acyl, alkyl, alkenyl, alkynyl, alkoxy, cycloalkyl; R3 is H, nitro, CN, N3, NH2, halogen, sulfonyl-amino, sulfenyl-amino, sulfenyl-amino, acyl, alkyl, alkenyl, alkynyl, cycloalkyl, alkoxy, alkylthio, alkylamino, alkyl-sulfinyl, alkyl-sulfonyl, alkyl-sulfonyl, R4-R6 are independently H, nalogen, CN, N3, NO2, alkyl, cycloalkyl, halo-alkyl, alkoxy, alkylthio, alkyl-sulfonyl, alkyl-sulfonyl, halo-alkyl, alkoxy, alkylthio, alkenyl, alkynyl, alkoxycarbonyl, amino, alkyl-sulfonyl, amino, alkyl-sulfonyl, amino, alkyl-sulfonyl, amino, alkyl-sulfonyl, amino, alkyl-sulfonyl, amino, alkylamino, amino-carbonyl, alkylaminocarbonyl, sulfonyl-amino, sulfenyl-amino, and acyl, which were prepared and tested as pesticides against animal pests for use in crop protection. Thus, I (R1 Me, R2 = i-Pr, R3 = OMe, R4 = R6 = H) was prepared from 3,5-dichloro-4-pyridinecarboxaldehyde by sequential nucleophilic substitution reactions with propane-1-thiol and with sodium methoxide, conversion of the thioether to a sulfonyl chloride with chlorine in chlorobenene, reaction of the sulfonyl chloride with holdrine in chlorobenene, reaction of the sulfonyl chloride with hydrogen sulfide. The title compds were used for combating animal pests from the orders Homoptera or Thysanoptera and were applied in an amount of 100 mg to 10 Kg per 100 Kg of seeds.

\*\*ECL-\*\*C-\*\*C-\*\*P-2-\*\*I-JP\*\*

\*\*EL RCT (Reactant): 59M (Synthetic preparation); PREP (Preparation); RACT (Reactant) of aminothiocarbonyl-substituted benzenesulfonamides and pyridinesulfonamides as pesticides for the protection of plants against animal pests and insects and particularly against aphids and thrips) \$50184-52-44 CABLUS

Benzenesulfonamide, 2-cyano-3-(difluoromethoxyl-N-2-propyn-1-yl- (CA INDEX NAME)

889097-44-3 CAPLUS Benzenesulfonamide, 2-cyano-N-ethyl-6-fluoro-3-methoxy-N-methyl- (CA INDEX NAME)

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Acetamide, N-[(2-cyano-3-methoxyphenyl)sulfonyl)-N-methoxy- (CA INDEX

937722-01-5 CAPLUS Acetamide, N-[[2-cyano-3-(difluoromethoxy)phenyl]sul[onyl]-N-ethyl- (CA INDEX NAME)

937722-02-6 CAPLUS Acetamide, N-[{2-cyano-6-fluoro-3-methoxyphenyl}sulfonyl]-N-ethyl- (CA INDEX NAME)

CAPLUS

Benzenesulfonamide, 2-cyano-N-(diethoxymethylene)-3-(difluoromethoxy)-(CA INDEX NAME)

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937722-22-0 CAPLUS Ethanimidic acid. N-[[2-cyano-3-(difluoromethoxy)phenyl]sulfonyl]-, methyl ester (CA INDEX NAME)

ANSWER 4 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
2006:1007694 CAPLUS Full-text
145:350188
Preparation of cyanobenzenesulfonamide derivatives as seed-treatment insecticides
Von Deyn, Wolfgang; Kaiser, Florian; Pohlman, Matthias; Bastiaans, Henricus Maria Martinus; Baumann, Ernst, Rack, Michael; Anspaugh, Douglas D.; Van Tuyl Cotter, Henry; Culbertson, Deborah L.; Hofmann, Michael; Hicks, Carol IN Hicks, Carol Basf Aktiengesellschaft, Germany

so	PCT Int CODEN:	. Appl., PIXXD2	65p	р.												
DT	Patent				•											
LA	English															
FAN.	CNT 1					*										
	PATENT	NO,		KIN	D	DATE		i i	APPL	ICAT	ION :	NO.		Di	ATE	
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PI	WO 2006	100288		A2		2006	928	1	WO 2	006-	EP60	988		2	00603	323
	WO 2006	100288		A3		2007	0125									
	₩;	AE, AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BW.	BY,	BZ,	CA,	CH,
		CN, CO,	CR,	Cυ,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
		GE, GH,														
		KZ, LC,														
		MZ, NA,														
		SG, SK,	SL,	SM,	SY,	TJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	υs,	UZ,	VC,
		VN, YU,														
	RW:	AT, BE,														
		IS, IT,														
		CF, CG,														
		GM, KE,					SD,	SL,	SZ,	Т2,	UG,	ZM,	ZW,	AM,	ΑŻ,	BY,
		KG, KZ,	MD,	RU,	ΤĴ,	TM										

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RI, AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses) (preparation as seed-treatment insecticide) 850154-03-79 CAPLUS

Benzenesulfonamide, 2-cvano-3-methoxy-N-methyl- (CA INDEX NAME)

850154-43-7 CAPLUS Benzenesulfonamide, 2-cyano-N-ethyl-3-methoxy- (CA INDEX NAME)

850154-45-9 CAPLUS Benzenesulfonamide, 2-cyano-3-methoxy-N-(1-methylethyl)- (CA INDEX NAME)

850154-46-0 CAPLUS Benzenesulfonamide, 2-cyano-3-methoxy-N-2-propynyl- (9CI) (CA INDEX NAME)

The cyanobenzenesulfonamide derivs. I [R1 = (halo)alkyl or (halo)alkoxy); R2 = H. (un)substitutec alkyl, alkomyl, alkynyl, cycloalkyl, etc.; R3-5 = H. halo. CN, NO2 (halo)alkyl, alkoxy, etc.] or its enantiomers are prepared as geed-treatment insecticides.
850154-03-P9 850154-47-IP 850154-48-2P
850154-96-3P 850154-47-IP 850154-68-2P
850154-96-3P 850154-61-P9 850154-68-2P
850154-96-3P 850154-61-P9 850154-68-2P
850154-96-P8 850154-69-PP 850154-70-0P
950154-71-IP 850154-72-2P 850154-70-0P
950154-71-SP 850154-72-2P 850154-70-0P
850154-80-5P 850154-80-9P 850154-80-0P
850154-96-P8 850154-98-9P 850154-80-0P
850154-96-P8 850154-90-4P 850155-98-0P
850154-96-P8 850154-90-4P 850155-98-0P
850154-96-P8 850154-96-0P 850155-07-4P
850155-09-P8 850155-07-5P 850155-07-4P
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850155-09-P8 850155-98-5P 850155-98-0P
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910459-47-IP 910459-85-3P 910459-46-0P
910459-47-IP 910459-85-3P 910459-67-9P
910459-57-3P 910459-85-3P 910459-67-9P
910459-67-97 910459-85-3P 910459-67-9P
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910459-97-3P 910459-78-3P 910459-77-PP
910459-97-3P 910459-78-3P 910459-78-7P
910459-97-3P 910459-78-3P 910459-88-0P

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850154-47-1 CAPLUS Benzenesulfonamide, 2-cyano-N-(cyanomethyl)-3-methoxy- (CA INDEX NAME)

850154-48-2 CAPLUS Benzenesulfonamide, 2-cyano-3-methoxy-N-2-propenyl- (9CI) (CA INDEX NAME)

850154-49-3 CAPLUS Benzenesulfonamide, 2-cyano-3-methoxy- (CA INDEX NAME)

850154-52-8 CAPLUS Benzenesulfonamide, 2-cyano-3-ethoxy-N-methyl- (CA INDEX NAME)

850154-53-9 CAPLUS

Benzenesulfonamide, 2-cyano-3-ethoxy-N-ethyl- (CA INDEX NAME)

850154-61-9 CAPLUS Benzenesulfonamide, 2-cyano-3-ethoxy-N-2-propynyl- (9CI) (CA INDEX NAME)

850154-62-0 CAPLUS Benzenesulfonamide, 2-cyano-N-(cyanomethyl)-3-ethoxy- (CA INDEX NAME)

850154-63-1 CAPLUS Benzenesulfonamide, 2-cyano-3-ethoxy-N-2-propenyl- (9CI) (CA INDEX NAME)

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850154-70-0 CAPLUS
Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-ethyl- (CA INDEX NAME)

850154-71-1 CAPLUS Benzenesulfonamide, 2-cyano-3-(1-methylethoxy)- (CA INDEX NAME)

850154-72-2 CAPLUS Benzenesulfonamide, 2-cyano-3-(1-methylethoxy)-N-2-propynyl- (9CI) (CA INDEX NAME)

850154-73-3 CAPLUS Benzenesulfonamide, 2-cyano-N-(cyanomethyl)-3-(1-methylethoxy)- (CA INDEX NAME)

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850154-64-2 CAPLUS Benzenesulfonamide, 2-cyano-3-ethoxy-N-propyl- (CA INDEX NAME)

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850154-67-5 CAPLUS Benzenesulfonamide, 2-cyano-3-ethoxy-N-(2,2,2-trifluoroethyl)- (CA INDEX NAME)

850154-68-6 CAPLUS Benzenesulionamide, 2-cyano-N-ethyl-3-(2-propenyloxy)- (9CI) (CA INDEX NAME)

850154-69-7 CAPLUS Benzenesulfonamide, 2-cyano-N-ethyl-3-(1-methylethoxy)- (CA INDEX NAME)

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850154-75-5 CAPLUS Benzenesulfonamide, 2-cyano-3-(1-methylethoxy)-N-(1-methylethyl)- (CA INDEX NAME)

850154-79-9 CAPLUS Benzenesulfonamide, 2-cyano-N-ethoxy-3-(1-methylethoxy)- (CA INDEX NAME)

850154-80-2 CAPLUS Benzenesulfonamide, 2-cyano-3-(1-methylethoxy)-N-propyl- (CA INDEX NAME)

RN 850154-81-3 CAPLUS
CN Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)- (CA INDEX NAME)

RN 850154-82-4 CAPLUS
CN Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-2-propyn-1-yl- (CA
INDEX NAME)

RN 850154-83-5 CAPLUS CN Benzenesulfonamide, 2-cyano-3-(2,3-dichloropropoxy)- (CA INDEX NAME)

RN 850154-84-6 CAPLUS CN Benzenesulfonamide, 2-cyano-N-methyl-3-(1-methylethoxy)- (CA INDEX NAME)

RN 850154-87-9 CAPLUS CN Benzenesulfonamide, 2-cyano-N-ethoxy-3-methoxy- (CA INDEX NAME)

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RN 850154-92-6 CAPLUS
CN Benzenesulfonamide, 2-cyano-3-methoxy-N-(2,2,2-trifluoroethyl)- (CA INDEX NAME)

RN 850154-93-7 CAPLUS
CN Bonzenesulfonamide, 2-cyano-N-(2,2-difluoroethyl)-3-methoxy- (CA INDEX NAME)

RN 850154-94-8 CAPLUS CN Benzenesulfonamide, 2-cyano-N,3-dimethoxy- (CA INDEX NAME)

RN 650154-95-9 CAPLUS
CN Benzenesulfonamide, 4-bromo-2-cyano-3-methoxy-N-2-propynyl- (9CI) (CA INDEX NAME)

N 850154-88-0 CAPLUS N Benzenesulfonamide, 2-cyano-N-{2-cyanoethyl}-3-methoxy- (CA INDEX NAME)

RN 850154-89-1 CAPLUS
CN Benzenesulfonamide, 2-cyano-3-methoxy-N-[2-(methylthio)ethyl]- (CA INDEX NAME)

RN 850154-90-4 CAPLUS
CN Benzenesulfonamide, 2-cyano-3-methoxy-N-{2-(methylsulfonyl)ethyl}- (CA INDEX NAME)

RN 850154-91-5 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-(2-fluoroethyl)-3-methoxy- (CA INDEX NAME)

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RN 850154-96-0 CAPLUS
CN Benzensulfonamide, 2-cyano-N-{2-{dimethylamino}ethyl}-3-methoxy- (CA INDEX NAME)

RN 850154-98-2 CAPLUS
CN Benzensulfonamide, 2-cyano-3-(difluoromethoxy)-N-(2,2,2-trifluoroethyl)(CA INDEX NAME)

RN 850155-01-0 CAPLUS
CN Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-[2-(methylthio)ethyl](CA INDEX NAME)

RN 850155-03-2 CAPLUS
CN Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-2-propenyl- (9CI) (CA

\$50155-05-4 CAPLUS Renzenesulfonamide, 2-cyano-3-(ditluorometnoxy)-N-propyl- (CA INDEX NAME)

850155-06-5 CAPLUS Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-(2-methoxyethyl)- (CA HOEX NAME)

0== S-NH-CH2-CH2-OME

850155-07-6 CAPLUS
Benzenesultonamide, 2-cyano-N-(2-cyanoethyl)-3-(difluoromethoxy)- (CA
INDEX NAME)

850155-08-7 CAPLUS

10574153

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850155-97-4 CAPLUS Benzenesul(onamide, 2-cyano-4-methoxy-3-methyl-N-2-propynyl- (9CI) (CA INDEX NAME)

850155-98-5 CAPLUS Benzenesulfonamide, 2-cyano-4-methoxy-3-methyl- (CA INDEX NAME)

850155-99-6 CAPLUS Benzenesulfonamide, 2-cyano-4-methoxy-N,3-dimethyl- (CA INDEX NAME)

910459-14 2 CAPLUS Benzenesulfonamide, 2-cyano-3-{difluoromethoxy}-N-methyl- (CA INDEX NAME)

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CN Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-(1-methylethyl)- (CA INDEX NAME)

F2CH- O CN CN O MHPr-1

850155-09-8 CAPLUS
Benzeneaulfonamide, 2-cyano-N-(2,2-difluoroethyl)-3-(difluoromethoxy)-(CA INDEX NAME)

S- NH- CH 2- CHF 2

850155-10-1 CAPLUS Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-methoxy- (CA INDEX NAME)

\$50155-96-3 CAPLUS Benzenesulfonamide, 2-cyano-N-ethyl-4-methoxy-3-methyl- (CA INDEX NAME)

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910459-15-3 CAPLUS Benzenesulfonamide, 4-bromo-2-cyano-3-methoxy-N-methyl- (CA INDEX NAME)

910459-31-3 CAPLUS Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-methyl- (CA INDEX NAME)

910459-45-9 CAPLUS Benzenesulfonamide, 2-cyano-N-ethyl-3-(trifluoromethoxy)- (CA INDEX NAME)

RN 910459-46-0 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-(1-methylethyl)-3-(trifluoromethoxy)- (CA INDEX NAME)

RN 910459-47-1 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-2-propynyl-3-(trifluoromethoxy)- (9CI) (CA INDEX NAME)

RN 910459-49-3 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-(2-methoxyethyl)-3-(trifluoromethoxy)- (CA INDEX NAME)

RN 910459-50-6 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-[2-(methylthio)ethyl]-3-(trifluoromethoxy)(CA INDEX NAME)

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CN Benzenesulfonamide, 2-cyano-3-(dichlorofluoromethoxy)-N-ethyl- (CA INDEX NAME)

RN 910459-56-2 CAPLUS
CN Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-(1-methylethyl)(CA INDEX MAME)

RN 910459-57-3 CAPLUS
CN Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-2-propynyl- (9CI)
(CA INDEX NAME)

RN 910459-59-5 CAPLUS
CN Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-(2-methoxyethyl)(CA INDEX NAME)

O S NH - CH2 - CH2 - SM
CN
O - CP3

RN 910459-51-7 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-propyl-3-(trifluoromethoxy)- (CA INDEX NAME)

RN 910459-52-8 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-(2,2,2-trifluoroethyl)-3-(trifluoromethoxy)(CA INDEX NAME)

RN 910459-53-9 CAPLUS
CN Benzenesulfonamide, 2-cyano-N-2-propenyl-3-(trifluoromethoxy)- (9CI) (CA INDEX NAME)

RN 910459-55-1 CAPLUS

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RN 910459-60-8 CAPLUS
CN Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-[2-(methylthiolethyll- (CA INDEX NAME)

RN 910459-61-9 CAPLUS CN Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-propyl- (CA INDEX NAME)

RN 910459-62-0 CAPLUS
CN Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-(2,2,2-trifluoroethyl)- (CA INDEX NAME)

RN 910459-63-1 CAPLUS
CN Banzaneaulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-2-propenyl- (9C1)
(CA INDEX NAME)

910459-66-4 CAPLUS
Benzenesulfonanide, 3-(chlorodifluoromethoxy)-2-cyano-N-[1-methyl-2-methylniolethyl]- (CA INDEX NAME)

910459-67-5 CAPLUS Benzenesultonamide, 3-(chlorodifluoromethoxy)-2-cyano-N-(2-cyanoethyl)-(CA INDEX NAME)

5- NH- CH2-- CH2-- CN

910459:69 7 CAPLUS
Bentzenesulfonamidu, 2-cyano-N-[1-methyl-2-(methyltnio)ethyl]-5-ttrifluoromethoxy; (CA INDEX NAME)

910459-71-1 CAPLUS Benzenesultonamide, 2-cyano-N-(2-cyanoethyl)-3-(trifluoromethoxy)- (CA INDEX NAME)

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910459-20-2 CAPLUS Benzenesulfonamide, 2-cyano-4-fluoro-3-methoxy-N-methyl- (CA INDEX NAME)

910459-81-3 CAPLUS Benzenesulfonamide, 2-cyano-4-fluoro-3-methoxy-N-(1-methylethyl)- (CA INDEX NAME)

910459-92-4 CAPLUS Benzenesulfonamide, 2-cyano-4-(luoro-3-methoxy-N-2-propynyl- (9CI) (CA

910459-83 5 CAPLUS

\_ ў— NН— СН 2— СН2 — СМ

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910459-73-3 CAPLUS Benzenesulfonamide, 2-cyano-4-(difluoromethoxy)-N-ethyl-3-methyl- (CA INDEX NAME)

910459-74-4 CAPLUS
Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N-(3,4,4-crifluoro-3-butenyl)-(9C1) (CA INDEX NAME)

O NH CH2- CH2- C- F

910459-77-7 CAPLUS Benzenesulfonamide, 2-cyano-N-ethyl-3-methoxy-4-methyl- (CA INDEX NAME)

910459-78-8 CAPLUS Benzenesulfonamide, 2-cyano-3-methoxy-4-methyl-N-2-propynyl- (9CI) (CA INDEX NAME)

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Benzenesulfonamide, 2-cyano-N-ethyl-4-fluoro-3-methoxy- (CA INDEX NAME)

910459-94-6 CAPLUS Benzenesulfonamide, 2-cyano-4-fluoro-3-methoxy-N-[2-(methylchio)ethyl]-(CA INDEX NAME)

L22 ANSMER S OF 15 CAPLUS COPYRIGHT 2008 ACS ON STN

AN 2006:1005623 CAPLUS Pull-text

DN 145:329902

1 Preparation of sulfonyl insecticides for seed treatment

IN Yon Deyn, Wolfgang; Kaiser, Florien; Pohlaan, Matthias; Bastiaans,
Henricus Maria Martinus; Baumann, Ernst; Rack, Michael; Anspaugh, Douglas

D.; Cotter, Henry Van Tuyl; Culbertson, Deborah L.; Hofmann, Michael;
Hicks, Carol

PA Basf Aktiengesellschaft, Germany

PO PCT Int. Appl., 44pp.

CODEN: PIXXD2

P Patent

LA English
FAN.CNT I

PATENT NO. KIND DATE APPLICATION NO. DATE

EP 1863349 A1 20071212 EP 2006-725244 20060322 AT 1853-949 A1 20071212 EP 2006-725244 20060322
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, CB, GR, HU, IE,
IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR
PRAI US 2005-6647173P P 20050324
NO 2006-EP60961 W 200660322

WO 2006-EP60961 MARPAT 145:329902

The sulfonyl compds. I (RI = halo; R2 = H, (halo)alkyl, (halo)alkenyl, (halo)alkynyl, etc.; R3-5 = H, halo, CN, NO2, (halo)alkyl, cycloalkyl, etc.) and their enantiomers are prepared as insecticides for seed treatment. 9c+904-05-89 304904-05-29 (halo)alkyl, etc.) RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (USeS) (preparation as insecticide for seed treatment) 909904-05-8 CAPLUS Benzenesulfonamide, 3-chloro-2-cyano-4-(difluoromethoxy)-N-ethyl- (CA INDEX NAME)

-06-9 CAPLUS esulfonamide, 3-chloro-2-cyano-N-ethyl-4-methoxy- (CA INDEX NAME)

909904-09-2 CAPLUS

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S-2,4,5-triaryl-imidazolines I, wherein X1 is alkoxy, X2 and X3 are independently H, halogen, CN, alkyl, alkoxy, piperidinyl, substituted amine, aulfonamide, acyl, sulfoxide, oxime, alkynyl, X2 and X3 together for heterocycle; Y1 and Y2 are independently H, acetylene; R is alkoxy, piperidinyl substituted with S-6 membered heterocycle or OH or COMM2, piperazinyl substituted by R1, R1 is H, Oxo, alkyl, acyl, sulfonyl, were prepared and used as antitumor agents. These compds. are believed to inhibit MPM2-p53 interaction and as such the compds. will have anti-hyperproliferative cellular activity. Thus, 5-{(45,5R)-4,5-bis(4-Chlorophenyl)-1-[4-(2-chloro-4-ethoxy-N,N-dimethylbenzenesulfonamide was prepared and used for the treatment of breast, colon, lung and prostate tumors. The ability of the compds. to inhibit in vitro the interaction between p53 and MDM2 proteins was measured. IC50s showing biol. activity that applies to compds. of the subject matter of this invention ranges from about 0.005 µM to about 1 µM, 5-[4,5-6] sig(4-chlorophenyl)-1-[4-(2-hydroxyethyl)-3-oxopiperazine-1-carbonyl]-4,5-dihydro-1H-imidazol-2-yl)-2-chloro-4-ethoxy-N,N-dimethylbenzenesulfonamide.

9:105/3-45-59 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation of cis-2,4,5-triaryl-imidazolines and their use as anticancer medicaments) 9:10568-65-9 CAPLUS Benzoic acid, 4-cyano-5-[(dimethylamino)sulfonyl}-2-ethoxy-, ethyl ester (CA INDEX NAME)

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 7 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN 2006:510657 CAPLUS Full-text 145:27730 Preparation of 2-cyano-3-(halo)alkoxybenzenesulfonamides as pesticides. Von Deyn, Molfgang, Bastiaans, Henricus María Martinus, Pohlman, Matchios, Rack, Michael; Baumann, Ernst; Pull, Michael, Hofmann, Michael; Todeschi, Livio; Kordes, Markus, Koradin, Christopher, Anspaugh, Douglas D.,

10574153

34 of 52 Benzenesulfonamide, 3-chloro-2-cyano-4-methoxy-N-2-propynyl- (9CI) (CA

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT RE.CNT 6

ANSWER 6 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN 2006:99005) CAPLUS FUll-text 145:377334

Preparation of cis-2.4,5-triaryl-imidazolines and their use as anticancer medicaments
Futouhi, Nader, Haley, Gregory, Jay, Simonsen, Klaus, B.; Vu, Binh, Thanh; Webber, Stephen, Evan F.Hoffmann-La Roche AG, Switz.
PCT Int. Appl., 293pp.
CODEN: PIXXD2
Patent IN PA SO DT Patent LA English FAN.CNT 1 PATENT NO. APPLICATION NO. KIND DATE 20060921 PI WO 2006097261 A1 20060921 WO 2006-EP2282 20060313

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, CE, GI, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MM, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SS, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, DA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, CF, CG, C1, CM, GA, GN, GO, CW, ML, MR, NE, SN, TD. TG, EW, GH, GK, KZ, MD, RU, TJ, TM

AU 2006224765 A1 20060921 AU 2006-224765 20060913

EP 1861376 A1 20060921 CA 2006-2299476 20060913

EP 1861376 A1 20060921 DS 2006-2374407 20060913

EP 1861376 A1 20060921 DS 2006-274765 20060913

ER, AT, BE, BG, CH, CY, CZ, DE, DK, EE, SF, FI, FR, GB, GR, HU, IE, KR 2007107129 A 20071123 IN 2007-CN4051 20070914

FRAI US 2005-662516F P 20050913

SMARPAT 145:377334

GI A1 WO 2006-EP2282 WO 2006097261 20060313

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Culbertson, Deborah L.; Cotter, Henry Van Tuyl; Oloumi-Sadeghi, Hassan Basf Aktiengesellschaft, Germany PCT Int. Appl., 44 pp. CODEN: PIXXD2 50 DT Patent LA English FAN.CNT 1 DT

PATENT NO KIND DATE APPLICATION NO. DATE CN 2005-80040653 IN 2007-KN1847 KR 2007-714451 CN 101065353 20071031 20051124 IN 2007KN01847 20070810 RR 2007086639
PRAI US 2004-631204P
WO 2005-EP12561
OS MARPAT 145:27730 20070827 20070625 20041126 20051124

Title compds. [I; A = alkoxy, haloalkoxy, Rl = alkyl, alkenyl; R2 = [substituted] alkyl, alkenyl, alkynyl, cycloalkyl, alkoxy, R3-R5 = H, halo, cyano, NO2, alkyl, cycloalkyl, haloalkyl, alkoxy, alkylthio, alkylsulfinyl, alkylsulfinyl, alkoxylandonyl, etc.], were prepared Thus, 2-cyano-3-difluoromethoxyphenylsulfonyl chloride (preparation given) and dimethylamine were stirred in THF/aqueous Na2CO3 to give 79% N,N-di-Me 2-cyano-3-difluoromethoxyphenylsulfonyl can the latter and addnl. I at 300 ppm gave >s5% kill of Aphis gossypii on cotton plants.

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(preparation of 2-cyano-3-(halo)alkoxybenzenesulfonamides as pesticides) 899097-30-7 CAPLUS Benzenesulfonamide, 2-cyano-3-(difluoromethoxy)-N,N-dimethyl- (CA INDEX NAME)

889097-31-8 CAPLUS Benzenesulfonamide, 2-cyano-3-methoxy-N,N-dimethyl- (CA INDEX NAME)

939097 32-9 CAPLUS Benzenesulfonamide, 2-cyano-N,N-diethyl-3-methoxy- (CA INDEX NAME)

859097-J3-0 CAPLUS Benzenesulfonamide, 2-cyano-3-methoxy-N,N-di-2-propenyl- (9CI) (CA INDEX NAME)

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889097-38-5 CAPLUS
Benzenesulfonamide, 2-cyano-3-methoxy-N,N,6-trimethyl- (CA INDEX NAME)

s39097-39-6 CAPLUS Benzenesulfonamide, 2-cyano-6-ethyl-2-methoxy-N,M-dimethyl- (CA INDEX NAME)

839097-40-9 CAPLUS
Bensoic acid, 3-cyano-2-[(dimethylamino)sulfonyl]-4-methoxy-, methyl ester
(CA INDEX NAME)

889097-41-0 CAPLUS Benzenesulfonamide. 6-chloro-2-cyano-3-methoxy-N,N-dimethyl- (CA INDEX NAME)

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889097-34-1 CAPLUS Benzenesulfonamide, 4-bromo-2-cyano-3-methoxy-N,N-dimethyl- (CA INDEX NAME)

889097-35-2 CAPLUS Benzenesulfonamide, 2-cyano-3-ethoxy-N,N-dimethyl- (CA INDEX NAME)

889097-36-3 CAPLUS Benzenesulfonamide, 3-(chlorodifluoromethoxy)-2-cyano-N,N-dimethyl- (CA INDEX NAME)

889097-37-4 CAPLUS
Benzenesulfonamide, 2-cyano-N,N-dimethyl-3-(trifluoromethoxy)- (CA INDEX
NAME)

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889097-42-1 CAPLUS
Benzenesulfonamide, 3-(2-chloro-1,1,2-trifluoroethoxy)-2-cyano-N-ethyl-N-methyl- (CA INDEX NAME)

889097-43-2 CAPLUS Benzenesulfonamide, 2-cyano-6-fluoro-3-methoxy-11,N-dimethyl- (CA INDEX NAME)

889097-44-3 CAPLU9 Benzenesulfonamide, 2-cyano-N-ethyl-6-fluoro-3-methoxy-N-methyl- (CA INDEX NAME)

889097-45-4 CAPLUS
Benzensuifonaside, 2-cyano-6-fluoro-3-methoxy-N-methyl-N-(1-methylethyl)(CA INDEX NAME)

859097-46-5 CAPLUS
BENZENESUITOnamide, 2-cyano-6-fluoro-3-methoxy-N-methyl-N-2-propynyl(9C1) (CA INDEX NAME)

889097-47-6 CAPLUS Benzenesulfonamide, 2-cyano-3,4-dimethoxy-N,N-dimethyl- (CA INDEX NAME)

889097-48-7 CAPLUS Senzenesulfonamide, 2-cyano-3-(difluoromethoxy)-6-fluoro-N,N-dimethyl-(CA INDEX NAME)

889097-49-8 CAPLUS Benzenesulfonamide, 6-bromo-4-chloro-2-cyano-3-methoxy-N,N-dimethyl- (CA INDEX NAME)

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The title compds. I [A is optionally substituted 3-sulfamoylphenyl, optionally substituted 2-thienyl, etc.; R10 is hydrogen or optionally substituted lower alkyl; R11 is hydroyen, halogeno, or hydroxy; R12 is hydrogen, halogeno, or nitro; R13 is hydroxy, optionally substituted lower alkoxy, optionally substituted lower alkoxy, optionally substituted lower alkoxy, optionally substituted lower alkylamino, etc.; R14 is hydrogen, halogeno, nitro, etc.; and R15 is hydrogen, halogeno, or hydroxy] are prepared In an in vitro binding assay for corticotropin releasing factor receptor antagonism, the potassium salt of title compound II showed IC50 of 36 nM.
357232-15-3P 357332-22-EP
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of benzamide derivs, as corticotropin releasing factor pror

receptor

ptor antagonists) 357332-59-3 CAPLUS Benzenesulfonamide, 3-cyano-N,N-diethyl-4-methoxy-5-nitro- (CA INDEX NAME)

357332-93-5 CAPLUS Benzonesulfonamide, 3-amino-5-cyano-N,N-diethyl-4-methoxy- (CA INDEX

RE.CNT 97

THERE ARE 97 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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889097-50-1 CAPLUS Benzenesulfonamide, 2-cyano-4-fluoro-3-methoxy-N,N-dimethyl- (CA INDEX

42 of 52

L22 ANSMER 8 OF 15 CAPLUS COPYRIGHT 2008 ACS ON STN
AN 2001:636036 CAPLUS <u>Full-text</u>
D1 135:210839
TI Preparation of benzamide derivatives as corticotropin releasing factor Dentation of the Management of the Miyazaki, Akira Japan Tobacco, Inc., Japan PCT Int. Appl., 221 pp. CODEN: PIXXD2

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рT Patent

LA	Japanes CNT 1	e								
· ru•	PATENT	NO.	KI	ND D	ATE	APPLI	CATION N	NO.	DATE	
PΙ	WO 2001	062718	А	1 2	0010830	WO 20	01-JP142	29	20010	226
	W:	AE, AG,	AL, AM	, AU,	AZ, BA,	BB, BG,	BR, BY,	BZ, CA,	CN, CR,	CU.
		CZ, DM,	DZ, EE	, GD, 0	GE, HR,	HU, ID,	IL, IN,	IS, KG,	KR, KZ,	LC.
		LK, LR,	LT, LV	, MA, I	MD, MG,	MK, MN,	MX, NO,	NZ, PL,	RO, RU,	SG.
		SI, SK,	TJ, TM	, TT, 8	UA, US,	UZ, VN,	YU, ZA,	AM, AZ,	BY, KG,	ΚZ,
		MD, RU,	TJ, TM							
	RW:	GH, GM,	KE, LS	, MW, I	MZ, SD,	SL, SZ,	TZ, UG,	ZW. AT,	BE, CH,	CY.
		DE, DK,	ES, FI	, FR, 0	GB, GR,	IE, IT,	LU, MC,	NL, PT.	SE, TR,	BF,
		BJ, CF,	CG, CI	, CM, 0	GA, GN,	GW, ML,	MR, NE,	SN, TD,	TG	
	JP 2002	201172	A	21	0020716	JP 20	01-45859	•	20010	221

JP 2002201172
PRAI JP 2000-50336
JP 2000-337197
JP 2000-337579
OS MARPAT 135:210839

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L22 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2008 ACS ON STN AN 1988:473125 CAPLUS <u>Full-text</u>
DN 109:73125

Preparation of phenoxyaminopropanols as β-blockers and diuretics Shioiri, Takayuki, Aoyama, Toyohiko, Takamori, Masayuki

Japan
Jpn. Kokai Tokkyo Koho, 10 pp.
CODEN: JKXXAF
Patent

LA Japanese FAN.CNT 1 PATENT NO.

MIND DATE
A 19870922
19860317 APPLICATION NO.
JP 1986-58982 PI JP 62215559 PRAI JP 1986-58982 OS CASREACT 100 T

CASREACT 109:73125

The title compds. I (R = alkyl; X = alkyl, sulfamoyl, halo, alkoxycarbonyl, The title compds: I (R = alkyl; X = alkyl, sulfamoyl, halo, alkoxycerbonyl, carbamoyl, cyano, alkoxy, alkoxycerbonylamino, atc.) were prepared as β-blockers and diuretics (no data). Etherification of II with CH2:CHCH2Br followed by oxidation with m-CloSH4CO2OH and reaction with Me2CHNH2 gave I,HC1 (R = Me2CH, X = Me).

115582-97-2P 115582-99-4P RE: SPN (Synthetic preparation); PREP (Preparation) (preparation of, as β-blocker and diuretic)

115582-97-3 CAPLUS

Benzenesulfonamido, 2-chloro-5-cyano-4-{2-hydroxy-3-[(1-methylethyl)amino]propoxy]-, monohydrochloride (9CI) (CA INDEX NAME)

115582-98-4 CAPLUS
Benzenesulfonamide, 2-chloro-5-cyano-4-[3-[(1,1-dimethylethyl)amino]-2-hydroxypropoxy]-, monohydrochloride (9CI) (CA INDEX NAME)

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45 of 52

● BCI

IT RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, in preparation of β-blocker and diuretic)
115522 Fd-4 CAPLUS
Benzenesulfonamide, 2-chloro-5-cyano-4-(2-propenyloxy)- (9CI) (CA INDEX

L22 ANSWER 10 OF 15 CAPLUS COPYRIGHT 2008 ACS ON STN
AN 1982:6713 CAPLUS Full-text
DN 96:6713
OREF 96:1222h,1223a
TI Sulfonyl compounds and aphicidal comnession
N van Hes, Roelof: Green Sulfonyl compounds and aphicidal compositions based on these compounds van Hes, Roelof; Grosscurt, Arnoldus Cornelis, Balk, Mouter Duphar International Research B. V., Neth. Eur. Pat. Appl., 48 pp. CODEN: EPAXEM

DT Patent LA English FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	• - •				
PI	EP 33984	A1	19810819	EP 1981-200042	19810115
	EP 33984	В1	19940606		
	R: AT, BE, CH,	DE, FR	, GE, IT, NI	., SE	
	AT 7787	T	19840615	AT 1981-200042	19810115
	BR 8100281	A	19810804	ER 1981-281	19810119
	US 4379157	A	19830405	US 1981-226533	19810119
	DK 8100241	A	19810724	DK 1981-241	19610120
	AU 9166360	A	19910730	AU 1981-66360	19810120
	JP 56120664	A	19810922	JP 1981-6043	19810120
	ZA 9100382	A	19920224	ZA 1981-382	19810120
	DD 157071	A5	19821013	DD 1981-227106	19810120
	RO 81451	A1	19840402	RO 1981-103171	19810120
	IL 61942	A	19951129	IL 1981-61942	19810120
	ES 498655	A1	19211101	ES 1981-498695	19810121

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The N-(1-alkyl-2-pyrrolidinylalkyl)benzamides I (R. R1 = alkyl, X = alkylene) or their acid addition salts were prepared by oxidation of their S analogs II or their acid addition salts. Thus, 1 g II (R = Me, R1 = Et, X = CH2), obtained from O-Me 2-methoxy-5-sultamoylthiobenzoate and 2-(aminomethyl)-1-ethylpyrrolidine, was refluxed with 2 g Pb(OAc)4 in H2O for 6 hr to give 0.75 g I (R = Me, R1 = Et, X = CH2). AB 17

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L22 ANSWER 12 OF 15 CAPLUS COPYRIGHT 2008 ACS ON STN AN 1975:4117 CAPLUS Fall-text DN 52:4117 OREF 62:7074,710a N-[1-Ethyl-2-pyrrolidinyl]methyl]-2 methoxy-5 sultamoylbenzamide Bulteau, Gerard; Acher, Jacques; Monier, Jean C. Societe d'Etudes Scientifiques et Industrielles de l'Ile-de-France

so

Ger. Otten., 9 pp. CODEN: GWXXBX

DТ Patent German

FAN. CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 2409891	A1	19740912	DE 1974-2409891	19740301
	FR 2220522	A1	19741004	FR 1973-8033	19730306
	FR 2220522	B1	19770204		
	CA 1010046	A1	19770510	CA 1974-193458	19740225
	AT 7401590	A	19790415	AT 1974-1590	19740227

10574153		46 of 52		
CS 219348	B2	19830325	CS 1981-439	19810121
CA 1167043	A1	19840508	CA 1991-368982	19810121
HU 28630	A2	19831228	HU 1981-134	19810122
HU 186379	В	19850729		
ES 500638	A1	19820101	ES 1981-500638	19810324
SU 1093247	A3	19840515	SU 1981-3328851	19810619
PRAI NL 1980-414	A	19800123		
EP 1981-200042	A	19810115		
OS MARPAT 96:6713				
GI				

$$R = \begin{bmatrix} R^1 & & & & \\ & & & & \\ & & & & \\ So_2R^2 & 1 & & & \\ \end{bmatrix} \begin{bmatrix} c_1 & & & \\ & & & \\ & & & \\ S_2^{**} & & \\ & & & \\ \end{bmatrix}$$

Sulfones I (R = halogen, OPh, alkyl, alkoxy; Rl = cyano, R2 = amino; R1R2 = \$,\$-dialkylsulfoximido, 1-amino-2-azavinylene, 1-amino-2-azaechylene) were prepared Thus 11.8 g 3,2-Cl(RC)C6H3802Cl was treated with 12.5 mL 25% aqueous NH3 Lo give 8.9 g II which was aphicidal at 10 mg/L. \$3001-65-EP RL: SPN (Synthetic preparation); PREP (Preparation) (preparation and aphicidal activity of) 80043-65-2 CAPLUS Benzenesulfonamide, 2-cyano-N,N-dimethyl-5-(trifluoromethoxy)- (CA INDEX NAME) AB

L22 ANSMER 11 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
AN 1977;5307 CAPLUS Full-text
DN 86:5307
EN 86:915a,918a
TI N-(1-81kyl-2-pyrrolidinylalkyl)-2-alkoxy-5-sulfamoylbenzamides
PA Societe d'Etudes Scientifique et Industrielles de l'Ile-de-France, Fr.
CODEN, JAXXAF
DT PATENT
LA Japanese
FARLCMT 1
PATENT NO. KIND DATE APPLICATION NO. DATE JP 1974-78617 19760123 19740709

L22 ANSMER 13 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
AN 1969:461019 CAPLUS <u>Full-text</u>
N 71:61019
OREF 71:11211a,11214a
T1 2-Alkowy-5-sulfamidobenzoic acids
Societe d'Etudes Scientifiques et Industrielles de l'Ile-de-France
CODEN: FRXXAK

DT LA

FAN.	CNT 1				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	FR 1524398		19680510	FR 1967-101151	19670331
АВ	solution of 200 g. thoxybenzenesulfony (d. 1.49) at -6° ga Reaction of 200 g. nitro-4-sulfamoylan by hydrogenation in g. Raney Ni, gave 2 of 81 g. IV, 424 ml	anisole i chlor ve 211 II with isole ( an aut 5 g. 2 water	in 700 ml. ide (I), m. g. 3-nitro-4 600 g. ammo III), m. 142 oclave at 50 methoxy-5-su , and 77 ml.	addition at -10° of 43 CHCl3 gave 210 g. 4-me- 38°. i (205 g.) added -methoxy- benzenesulfon nium carbonate at 100° °. III (30 g.) in 50 m (730 atmospheric in the lfamoylaniline (IV), m. HCl (d. 1.19) treated dat 30° to a solution	to 718 ml. HNO3 yl chloride (II). gave 172 g. 2- l. absolute EtOH, presence of 20 143°. A mixture at 0° with 32.3

Cus04.5H20 in 645 ml. water and then reacted with 89.2 g. NaCN gave 41 g. 2-methoxy-5-sulfamoylbenzonitrile (V), m. 180-1°. A mixture of 15 g. V, 74 ml. water, 70 ml. H2504 (d. 1.84) heated to 130-40° for 30 min. gave 11 g. 2-methoxy-5-sulfamoylbenzoic acid, m. 220°. Similarly, from 2-ethoxybenzene, 2-ethoxy-5-(dimethylsulfamoyl)-benzoic acid, m. 123°, was prepared 22117-34-6P 22121-74-0P

17 RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of) 22117-84-6 CAPLUS

Benzenesulfonamide, 3-cyano-4-methoxy- (CA INDEX NAME)

22121-74-0 CAPLUS
Benzenesulfonamide, 3-cyano-4-ethoxy-N,N-dimethyl- (CA INDEX NAME)

L22 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN AN 1969:430239 CAPLUS Full-text DN 71:30239

71:55259 71:5569a.5572a ORFE

71:5594.55/28
2-Alkoxy-5-sulfonamidobenzoic acids
Societe d'Etudes Scientifiques et Industrielles de l'Ile-de-France
Fr., 4 pp.
CODEN: FREXXAK

DT LA FAN

CODEN: PRXXAK
PATENT
PRINCH
French

CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

FR 1532332 19680712 FR 1967-100123 19670323

For diagram(s), see printed CA Issue.
In the title compdo. (1), R = C.1-5 alkyl or alkenyl and R1,R2 = C1-3 alkyl groups, or NR1R2 is part of a ring containing O or N. I were prepared by a series of reactions starting with ordinary phenols. E.g., 470 g. PhOH and 650 g. concentrated H2S04 were stirred at 125-30° 6 hrs., then after dilution with H2O poured into a solution of 535 g. NaNO3 in 1 l. H2O and worked up to give

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dried and triturated with alc., the 2 filtrates added to the main solution, and the solution cooled 1 hr. and evaporated in vacuo to give 86% 2.5(MeO) (H2NSO2) C6H3NH2 (II), m. 142°. Similarly prepared was 70% 2.5(EtO) (Me2NSO2) C6H3NH2, m. 125°. A solution of 32.3 g. NaNO2 in 10% g. H2O was added to a solution of 81 g. II, 77 ml. HCI (d. 1.19), and 424 ml. H2O and the mixture stirred 30 min. at 0.5° and added to a diazo solution (prepared from 89.2 g. NaNO2 10.72 g. CuSO4, and 645 ml. H2O) at 85-90° in 15 min. the mixture stirred 20 min. at 85-90°, cooled, kept at 0.5° 2 hrs., and filtered, the solid washed in water, dried at 60°, and mixed with 1650 ml. absolute EtOH, the mixture filtered, and the filtrate boiled 0.5 hr. with charcoal, evaporated to 600 ml., and cooled 1 hr. to give 48.5% 2.5° (MEO) (H2NSO2) C6H3CN (III), m. 180-1°. Similarly prepared was 46.5% 2.5° (MEO) (ME2NSO2) C6H3CN, m. 103-4°. A mixture of 15 g. III, 70 ml. H2SO4 (d. 1.84), and 7 and . H2O was kept at 130-40° 0.5 hrs., cooled at 0.5° 2 hrs., and filtered, the solid triturated 4 times with H2O. dried at 60°, and dissolved in 250 ml. saturated NaHCO3 solution, the solution clarified with 2.5 g. charcoal 0.5 hr. and acidified to pH 1 with 25 ml. HCI (d. 1.19), and the precipitate filtered off, washed with H2O until the washings were C1 free, and dried at 60° to give 67% (2.5° (MEO) (MEONSO2) C6H3CO2H, m. 122°. Similarly prepared was 2,5° (EtO) (MEONSO2) C6H3CO2H, m. 122°. Similarly prepared was 2,5° (EtO) (MEONSO2) C6H3CO2H, m. 122°. Similarly prepared was 2,5° (EtO) (MEONSO2) C6H3CO2H, m. 122°.

ELSPN (Synthetic preparation), PREP (Preparation) (preparation) (preparation) 3° (PREP (Preparation) (CA INDEX NAME)

22121-74-0 CAPLUS

Benzenesulfonamide, 3-cyano-4-ethoxy-N,N-dimethyl- (CA INDEX NAME)

-> log hold COST IN U.S. DOLLARS

FULL ESTIMATED COST

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500 152 SOURCE AND THE STREET OF THE STREET

IT RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of)
22117-84-6 CAPLUS
Benzenesulfonamide, 3-cyano-4-methoxy- (CA INDEX NAME) RN

L22 ANSWER 15 OF 15 CAPLUS COPYRIGHT 2008 ACS on STN
1969:67925 CAPLUS Full-text
DN 70:67925
CAPLUS Full-text
DN 70:67925
TI 2-Alkoxy-5-sulfamoylbenzoic acids
Bulleau, Gerard
Societe d'Etudes Scientifiques et Industrielles de l'Ile-de-France
S 5. African, 18 pp.
CODEN: SFXXAB

GB 1204406

Patent

LA English

FAN	. CN1 I				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΙ	ZA 6801840		19680821	ZA	
	FR 1524388			FR	
	FR 1525352			FR	
	FR 1532332			FR	

PRAI FR

TR 1923322 FR 19670323 FR 19670323 FR 19670324 FR 19670323 FR 19670323 FR 19670406

A mixture of 600 g. crushed (NH4) 2CO3 and 200 g. 2-methoxy-5-chlorosultonyInitrobenzene was stirred in a boiling water bath 2 hrs., poured onto 2 kg. water and ice, and stirred 15 min., the solid filtered off, repeatedly (until foams no longer form) triturated with 200 ml. 10% HCl and with water, and dried at 60° to give 172 g. 2,5-(MeO) (H2NSO2) CEHSNO2 (1). Similarly prepared was 97% 2,5-(ECO) (Me2NSO2) CEHSNO2, m. 100-9°. A mixture of 206 g. powdered Fe and 800 ml. H20 was added to a stirred mixture of 158 g. I and 800 ml. absolute EtOH at 60°, the mixture stirred at 80° to suspend the Fe, a solution of 20 ml. HCl (d. 1.19) in 80 ml. H20 added <09°, the mixture kept at 80-5° 3 hrs., the pH adjusted to 7.5 by addition of a solution of 20 g. potash in 50 ml. H20 and 50 ml. alc., the mixture filtered, the brown solid

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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SESSION - 21.60 ENTRY -12.00 CA SUBSCRIBER PRICE

SESSION WILL BE HELD FOR 120 MINUTES STN INTERNATIONAL SESSION SUSPENDED AT 13:05:20 ON 28 JAN 2008